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<u>L4</u>	(gp39 or cd40L or cd40 adj ligand or 5c8) same (antibod\$) and ('89-76' or '24-31')	23	<u>L4</u>
<u>L3</u>	(gp39 or cd40L or cd40 adj ligand or 5c8) same (antibod\$)same (diabetes)	29	<u>L3</u>
<u>L2</u> <u>L1</u>	(gp39 or cd40L or cd40 adj ligand or $5c8$) same (antibod\$) and diabetes noelle-randolph\$	305 18	<u>L2</u> <u>L1</u>

END OF SEARCH HISTORY

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Search Results - Record(s) 1 through 10 of 23 returned.

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Search Results - Record(s) 11 through 20 of 23 returned.

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Search Results - Record(s) 21 through 23 of 23 returned.

☐ 21. <u>5876718</u> . 27 Mar 98; 02 Mar 99. Methods of inducing T cell non-resp	onsiveness to
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Term	Documents
GP39	237
GP39S	0
CD40L	877
CD40LS	2
CD40	2267
CD40S	0
LIGAND	58523
LIGANDS	47955
5C8	98
5C8S	0
89-76	21
((GP39 OR CD40L OR CD40 ADJ LIGAND OR 5C8) SAME (ANTIBOD\$) AND ('89-76' OR '24-31')).USPT,PGPB.	23

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L1: Entry 1 of 18

File: PGPB

Dec 12, 2002

PGPUB-DOCUMENT-NUMBER: 20020187135

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020187135 A1

TITLE: METHODS FOR INDUCING ANTIGEN-SPECIFIC T CELL TOLERANCE

PUBLICATION-DATE: December 12, 2002

INVENTOR - INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
NOELLE, RANDOLPH J.	CORNISH	NH	US	
FOY, TERESA M.	LEBANON	NH	US	
DURIE, FIONA H.	LEBANON	NH	US	

APPL-NO: 09/ 164568 [PALM]
DATE FILED: October 1, 1998

CONTINUED PROSECUTION APPLICATION: This is a publication of a continued prosecution application (CPA) filed under 37 CFR 1.53(d).

RELATED-US-APPL-DATA:

Application 09/164568 is a continuation-of US application 08/232929, filed April 25, 1994, US Patent No. 5869049

Application 08/232929 is a continuation-in-part-of US application 08/116255, filed September 2, 1993, ABANDONED

INT-CL: [07] $\frac{A61}{39/00}$, $\frac{K}{361}$ $\frac{38/16}{K}$, $\frac{A01}{1/00}$ $\frac{N}{1/00}$, $\frac{63/00}{1/00}$, $\frac{A61}{1/00}$ $\frac{K}{100}$, $\frac{39/395}{1/00}$, $\frac{A61}{1/00}$, $\frac{K}{100}$, $\frac{39/42}{1/00}$, $\frac{A61}{1/00}$, $\frac{K}{100}$, $\frac{39/40}{1/00}$, $\frac{A61}{1/00}$, $\frac{K}{100}$, $\frac{A61}{1/00}$, $\frac{A61}{1/00}$, $\frac{K}{100}$, $\frac{A61}{1/00}$,

 $\begin{array}{l} \text{US-CL-PUBLISHED: } 424/93.71; \ 424/133.1, \ 424/143.1, \ 424/173.1, \ 424/134.1, \ 424/153.1, \\ 424/141.1, \ 424/154.1, \ 424/184.1, \ 424/192.1, \ 424/577, \ 424/578, \ 514/8, \ 514/885, \ 530/350, \\ 530/387.3, \ 530/388.1, \ 530/388.2, \ 530/388.22, \ 530/388.7, \ 530/388.73, \ 530/388.75, \\ \text{US-CL-CURRENT: } 424/93.71; \ 424/133.1, \ 424/134.1, \ 424/141.1, \ 424/143.1, \ 424/153.1, \\ 424/154.1, \ 424/173.1, \ 424/184.1, \ 424/192.1, \ 424/577, \ 424/578, \ 514/8, \ 514/885, \ 530/350, \\ \hline 530/387.3, \ 530/388.1, \ 530/388.2, \ 530/388.2, \ 530/388.2, \ 530/388.75 \\ \end{array}$

REPRESENTATIVE-FIGURES: NONE

ABSTRACT:

Methods for inducing antigen-specific T cell tolerance are disclosed. The methods involve contacting a T cell with: 1) a cell which presents antigen to the T cell, wherein a ligand on the cell interacts with a receptor on the surface of the T cell which mediates contact-dependent helper effector function; and 2) an antagonist of the receptor on the surface of the T cell which inhibits interaction of the ligand on the antigen presenting cell with the receptor on the T cell. In a preferred embodiment, the cell which presents antigen to the T cell is a B cell and the receptor on the surface of the T cell which mediates contact-dependent helper effector function is gp39. Preferably, the antagonist is an anti-gp39 antibody or a soluble gp39 ligand (e.g., soluble CD40). The methods of the invention can be used to induce T cell tolerance to a soluble antigen or to an allogeneic cell. The methods of the invention can also be used to induce tolerance in cases of bone marrow transplantation and other organ transplants and to inhibit graft-versus-host disease.

RELATED APPLICATIONS

[0001] This application is a Continuation-in-Part of U.S. patent application Ser. No. 08/116,255, filed Sep. 2, 1993, the contents of which are incorporated herein by reference.

Print

Search Results - Record(s) 11 through 18 of 18 returned.

☐ 11. <u>6312692</u> . 30 Apr 98; 06 Nov 01. Method of treating graft-versus-host disease with anti-GP39 antibodies and bone marrow cells. <u>Noelle; Randolph J.</u> , et al. 424/154.1; 424/130.1 424/141.1 424/143.1 424/144.1 424/153.1 424/173.1 424/520 424/577 424/93.7 424/93.7 435/332 435/334 435/343 435/343.1 435/343.2 435/346 530/387.1 530/388.1 530/388.2 530/388.2 530/388.7 530/388.7 530/388.75. A61K039/395 A61K035/28 C07K016/28.
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☐ 14. <u>5876718</u> . 27 Mar 98; 02 Mar 99. Methods of inducing T cell non-responsiveness to transplanted tissues and of treating graft-versus-host-disease with anti-gp39 antibodies. <u>Noelle; Randolph J.</u> , et al. 424/154.1; 424/130.1 424/133.1 424/134.1 424/141.1 424/143.1 424/193.1 435/326 435/332 435/334 435/343 435/343.1 530/388.2 530/388.22 530/388.7 530/388.73 530/388.75. A61K039/395 C07K016/28 C12N005/12.
☐ 15. <u>5869049</u> . 25 Apr 94; 09 Feb 99. Methods of inducing T cell unresponsiveness to bone marrow with gp39 antagonists. <u>Noelle; Randolph J.</u> , et al. 424/154.1; 424/130.1 424/134.1 424/143.1 424/144.1 424/173.1 424/233.1 514/12 514/2 514/8. A61K039/395 A61K037/02 A61K037/04.
☐ 16. <u>5833987</u> . 07 Jun 95; 10 Nov 98. Treatment of T cell mediated autoimmune disorders. <u>Noelle;</u> <u>Randolph J.</u> , et al. 424/154.1; 424/130.1 424/133.1 424/141.1 424/142.1 424/143.1 424/144.1 424/153.1 424/173.1. A61K039/395.
☐ 17. <u>5747037</u> . 07 Jun 95; 05 May 98. Anti-GP39 antibodies. <u>Noelle; Randolph J.</u> , et al. 424/154.1; 424/130.1 424/141.1 424/143.1 424/144.1 424/153.1 424/173.1 435/326 435/332 435/334 435/343 435/343.1 435/343.2 435/346 435/70.21 530/387.1 530/388.1 530/388.2 530/388.22 530/388.7 530/388.75. A01K039/395 C07K016/28 C12N005/12.
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Term	Documents
NOELLE-RANDOLPH\$	0
NOELLE-RANDOLPH	2
NOELLE-RANDOLPH-J	16
NOELLE-RANDOLPH\$.USPT,PGPB.	18
(NOELLE-RANDOLPH\$).USPT,PGPB.	18

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L3: Entry 16 of 29

File: USPT

Jun 25, 2002

DOCUMENT-IDENTIFIER: US 6410711 B1

** See image for Certificate of Correction **

TITLE: DNA encoding CD40 ligand, a cytokine that binds CD40

Detailed Description Text (129):

These data indicate that the interaction of CD40 with its ligand is the principal molecular interaction responsible for T cell contact dependent induction of B cell growth and differentiation to both antigen-specific antibody production and polyclonal Ig secretion. As such, these data suggest that antagonists of this interaction, by soluble CD40, CD40/Fc fusion protein and possibly soluble CD40-L (monomeric), will significantly interfere with development of antibody responses. Therefore clinical situations where CD40, CD40/Fc fusion proteins and soluble CD40-L are suitable include allergy, lupus, rheumatoid arthritis, insulin dependent diabetes mellitus, and any other diseases where autoimmune antibody or antigen/antibody complexes are responsible for clinical pathology of the disease. Moreover, membrane-bound CD40-L or oligomeric soluble CD40-L will be useful to stimulate B cell proliferation and antibody production. As such, these forms of CD40-L are most useful for vaccine adjuvants and as a stimulating agent for mAb secretion from hybridoma cells.

L3: Entry 28 of 29

File: USPT

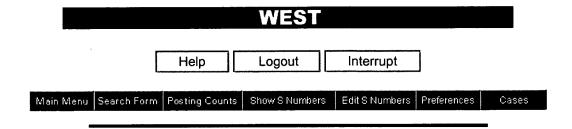
Mar 2, 1999

DOCUMENT-IDENTIFIER: US 5876950 A TITLE: Monoclonal antibodies specific for different epitopes of human GP39 and methods for their use in diagnosis and therapy

Detailed Description Text (30):
The pharmaceutical compositions of the present invention find use in vivo to inhibit the CD40/gp39 interaction. Blocking this interaction limits both primary and secondary antibody responses to T-cell dependent antigens and antibody production specific for these antigens. Therefore, the monoclonal antibodies, antigen binding fragments, and recombinant binding proteins can be used to inhibit the activation of B cells, modulating or inhibiting autoimmune disease (i.e., psoriasis, rheumatoid arthritis, systemic lupus erythematosis, diabetes mellitus, etc.), allergic responses, organ rejection or graft-versus-host disease. The compositions can also be used for imaging tumors which express gp39, when labeled with a detectable marker. When conjugated with

a therapeutic agent or as a fusion protein with a therapeutic agent, the monoclonal

antibodies, antigen binding fragment or recombinant binding proteins, can also be used o target the therapeutic agent to tumor cells.



Search Results -

Term	Documents
GP39	237
GP39S	0
CD40L	877
CD40LS	2
CD40	2267
CD40S	0
LIGAND	58523
LIGANDS	47955
5C8	98
5C8S	0
89-76	21
((GP39 OR CD40L OR CD40 ADJ LIGAND OR 5C8) SAME (ANTIBOD\$) AND ('89-76' OR '24-31')).USPT,PGPB.	23

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Term	Documents
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GP39S	0
CD40L	877
CD40LS	2
CD40	2267
CD40S	0
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LIGANDS	47955
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5C8S	0
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